WO 2005/056025 PCT/US2004/040178

20

## CLAIMS

- 1) A composition useful for the prevention, inhibition or treatment Parkinson's disease in a mammal comprising:
- a) live pigmented cells derived from the substantia nigra area of the brain of a mammal or the retinal pigmented epithelium layer or a mammal; and
- b) a biodegradable polymer gel capable of photo-induced cross linking.
- 2) The composition of claim 1 wherein said biodegradable polymer gel further comprises a water soluble macromer having poly(ethylene glycol) di-ethylphosphatidyl (ethylene glycol) methacrylate.
- 3) The composition of claim 2 wherein said biodegradable polymer gel further comprises attachment proteins and growth factors to enhance the survival of pigmented cells after implantation.
- 4) The composition of claim 3 wherein said attachment proteins can be laminin, fibronectin, and RGDS.
- 5) The composition of claim 1 wherein the live pigmented cells are mixed with the polymer gel solution (10 to 20% W/V).
- 6) The composition of claim 5 wherein the concentration of live pigmented cells is at least 200,000 cells to about 800,000 cells.

WO 2005/056025

21

PCT/US2004/040178

- 7) The composition of claim 3 wherein said growth factors are bFGF and EGF.
- 8) The composition of claim 7 wherein said growth factors are conjugated to polycarbophyll.
- 9) The composition of claim 1 wherein said biodegradable polymer gel further comprises a water soluble comprising a Poly-vinyl alcohol.
- 10) A method for the prevention, inhibition or treatment Parkinson's disease in a mammal comprising:
- a) harvesting pigmented cells (Human or bovine origin) from the brain stem (substantia nigra area) or from the retinal pigmented epithelium layer;
- b) maintaining said cells on BCE-ECM extracellular matrix coated dishes and suitable growth media;
  - c) harvesting at least 200,000 of said cells;
- d) preparing a mixture comprising a biodegradable polymer gel capable of photo-induced cross linking;
- e) mixing the live pigmented cells with the polymer gel solution (10 to 20% W/V);
- f) introducing into the brain of a mammal mixture of live pigmented cells with the polymer gel solution; and
- g) photo-polymerizing the polymer gel using UV light with a photoinitiator.
- 11) The method of claim 10, wherein said biodegradable polymer gel further comprises attachment proteins and growth factors to enhance the survival of pigmented cells after implantation.

- 12) The method of claim 10, wherein said attachment proteins can be laminin, fibronectin, and RGDS, and wherein said growth factors are bFGF and EGF.
- 13) The method of claim 10, wherein said biodegradable polymer gel further comprises a water soluble comprising a Poly-vinyl alcohol.
- 14) The method of claim 10, wherein said introduction into the brain of a mammal comprises injecting into the brain of a mammal the mixture of live pigmented cells with the polymer gel solution using a needle means.
- 15) A composition useful for the prevention, inhibition or treatment a retinal cell disease in a mammal comprising:
- a) live pigmented cells derived from the retina of a mammal; and
- b) a biodegradable polymer gel capable of photo-induced cross linking.
- 16) The composition of claim 15 wherein said biodegradable polymer gel further comprises a water soluble macromer having poly(ethylene glycol) di-ethylphosphatidyl (ethylene glycol) methacrylate.
- 17) The composition of claim 16 wherein said biodegradable polymer gel further comprises attachment proteins and growth factors to enhance the survival of pigmented cells after implantation.
- 18) The composition of claim 17 wherein said attachment proteins can be laminin, fibronectin, and RGDS.

WO 2005/056025 PCT/US2004/040178

- 19) The composition of claim 15 wherein the live pigmented cells are mixed with the polymer gel solution (10 to 20% W/V).
- 20) The composition of claim 19 wherein the concentration of live pigmented cells is at least 200,000 cells to about 800,000 cells.
- 21) The composition of claim 17 wherein said growth factors are bFGF and EGF.
- 22) The composition of claim 21 wherein the growth factors are conjugated to polycarbophyll.
- 23) The composition of claim 15 wherein said biodegradable polymer gel further comprises a water soluble comprising a Poly(vinyl alcohol).
- 24) A method for the prevention, inhibition or treatment of a retinal cell disease in a mammal comprising:
- a) harvesting pigmented cells (Human or bovine origin) from the retinal pigmented epithelium layer;
- b) maintaining said cells on BCE-ECM extracellular matrix coated dishes and suitable growth media;
  - c) harvesting at least 200,000 of said cells;
- d) preparing a mixture comprising a biodegradable polymer gel capable of photo-induced cross linking;
- e) mixing the live pigmented cells with the polymer gel solution (10 to 20% W/V);
- f) introducing into the retina of a mammal mixture of live pigmented cells with the polymer gel solution; and
  - g) photo-polymerizing the polymer gel using UV light

WO 2005/056025 PCT/US2004/040178

24

with a photoinitiator.

- 25) The method of claim 24, wherein said biodegradable polymer gel further comprises attachment proteins and growth factors to enhance the survival of pigmented cells after implantation.
- 26) The method of claim 24, wherein said attachment proteins can be laminin, fibronectin, and RGDS, and wherein said growth factors are bFGF and EGF.
- 27) The method of claim 24, wherein said biodegradable polymer gel further comprises a water soluble comprising a Poly (vinyl alcohol).
- 28) The method of claim 24, wherein said introduction into the retina of a mammal comprises injecting into the retina of a mammal the mixture of live pigmented cells with the polymer gel solution using a needle means.